MAR 2 7 2001

K010160

Section E

510(k) SUMMARY

Submitted by: Jensen Industries
50 Stillman Road
North Haven CT 06473
(203) 239-2090 phone
(203) 234-1015 fax
Contact: John Slanski

Date Prepared: January 15, 2001

Device Name: **Willi Geller Creapearl** denture teeth
Common Name: Synthetic Resin Denture Teeth
Classification: Class II
Product Code: ELM

Predicate Devices: K984128 – ENIGMA denture teeth (Davis Schottlander & Davis, Ltd.)
K984134 – NATURA denture teeth (Davis Schottlander & Davis, Ltd.)

Device Description:

Willi Geller Creapearl denture teeth are preformed, plastic denture teeth used in conjunction with a denture base resin for construction of removable full and partial dentures. The product is for use by dental professionals such as dentists and dental technicians. The line of denture teeth includes various shades, shapes, and sizes of anterior (ADA type I) and posterior (ADA type II) preformed plastic teeth.

Willi Geller Creapearl are certified by the manufacturer to comply with international standard ISO 3336 – Synthetic Polymer Teeth. This ensures safety and effectiveness of the device. ***Willi Geller Creapearl*** are manufactured from the same materials, in the same factory, as the predicate devices ENIGMA (K984128) and NATURA (K984134). For these reasons, ***Willi Geller Creapearl*** are substantially equivalent to the predicate devices.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAR 2 7 2001

Mr. John Slanski
Manager of Research & Development
Jensen Industries, Incorporated
50 stillman Road
North Haven, Connecticut 06473

Re: K010160
Trade Name: Willi Geller Creapearl
Regulatory Class: II
Product Code: ELM
Dated: January 15, 2001
Received: January 16, 2001

Dear Mr. Slanski:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does

not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsma/dsmamain.html".

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use Statement

Ver/ 3 - 4/24/96

Applicant: Jensen Industries Incorporated

510(k) Number (if known): K010160

Device Name: Willi Geller Creapearl

Indications For Use:

Willi Geller Creapearl teeth are to be used in conjunction with denture base resin for the construction of removable full and partial dentures.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)
(Per 21 CFR 801.109)
(Optional Format 1-2-96)

(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K010160